



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6: C07H 21/00, A61K 31/70	A3	(11) International Publication Number: WO 97/06662 (43) International Publication Date: 27 February 1997 (27.02.97)
(21) International Application Number: PCT/US96/13371 (22) International Filing Date: 16 August 1996 (16.08.96) (30) Priority Data: 08/516,454 17 August 1995 (17.08.95) US (71) Applicant: HYBRIDON, INC. [US/US]; One Innovation Drive, Worcester, MA 01605 (US). (72) Inventor: AGRAWAL, Sudhir; 61 Lamplighter Road, Shrewsbury, MA 01545 (US). (74) Agent: KEOWN, Wayne, A.; Hale and Dorr, 60 State Street, Boston, MA 02109 (US).		(81) Designated States: AL, AU, BB, BG, BR, CA, CN, CZ, EE, GE, HU, IL, IS, JP, KP, KR, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, TR, TT, UA, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i> (88) Date of publication of the international search report: 19 June 1997 (19.06.97)
(54) Title: INVERTED CHIMERIC AND HYBRID OLIGONUCLEOTIDES (57) Abstract The invention relates to modified oligonucleotides that are useful for studies of gene expression and for the antisense therapeutic approach. The invention provides inverted hybrid oligonucleotides and inverted chimeric oligonucleotides, both of which produce reduced side effects, relative to traditional phosphorothioate, hybrid or chimeric oligonucleotides.		

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A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C07H21/00 A61K31/70

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C07H A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	BIOORGANIC & MEDICINAL CHEMISTRY LETTERS, vol. 4, no. 24, 1994, pages 2929-2934, XP002020356 V. METELEV ET AL: "Study of Antisense Oligonucleotide Phosphorothioates Containing Segments of Oligodeoxyribonucleotides and 2'-O-Methyloligoribonucleotides" see the whole document ---	1-8
Y	WO 94 08003 A (ISIS PHARMACEUTICALS INC ;MONIA BRETT P (US); FREIER SUSAN M (US);) 14 April 1994 see page 26 - page 27 see page 30; table 6 --- -/--	1-8

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

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T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

28 April 1997

Date of mailing of the international search report

14.05.1997

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 268, no. 19, 5 July 1993, pages 14514-14522, XP000576145 MONIA B P ET AL: "EVALUATION OF 2'-MODIFIED OLIGONUCLEOTIDES CONTAINING 2'-DEOXY GAPS AS ANTISENSE INHIBITORS OF GENE EXPRESSION" see the whole document ---	1-8
Y	WO 95 02069 A (ISIS PHARMACEUTICALS INC ; BENNETT C FRANK (US); BOGGS RUSSELL T (U) 19 January 1995 see example 7 ---	1-8
Y	WO 93 13114 A (ISIS PHARMACEUTICALS INC) 8 July 1993 see examples 9,11,13 ---	1-8
Y	DE 41 10 085 A (BOEHRINGER INGELHEIM INT) 1 October 1992 see example 2 ---	1-8
X	WO 94 23028 A (HYBRIDON INC ;AGRAWAL SUDHIR (US); TANG JIN YAN (US); PADMAPRIYA A) 13 October 1994 ---	17-24
Y	see page 12 - page 15; examples A,B ---	17-24
Y	WO 94 02499 A (HYBRIDON INC ;PADMAPRIYA A (US); AGRAWAL SUDHIR (US)) 3 February 1994 see example 5 ---	17-24
Y	WO 91 12323 A (WORCESTER FOUNDATION FOR EX BI) 22 August 1991 see the whole document ---	17-24
Y	WO 94 07367 A (APOLLON INC ;MAX PLANCK GESSELLSCHAFT ZUR FO (DE); MOELLING KARIN () 14 April 1994 see page 13, paragraph 1; claims 27,28 ---	17-24
Y	ANTISENSE RESEARCH AND DEVELOPMENT, vol. 3, no. 1, 21 March 1993, pages 53-66, XP000611293 ZHAO Q ET AL: "COMPARISON OF CELLULAR BINDING AND UPTAKE OF ANTISENSE PHOSPHODIESTER, PHOSPHOROTHIOATE, AND MIXED PHOSPHOROTHIOATE AND METHYLPHOSPHONATE OLIGONUCLEOTIDES" see page 54, paragraph 2 ---	17-24

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>NUCLEIC ACIDS RESEARCH, vol. 19, no. 20, 1991, pages 5743-5748, XP002014768 HOKE G D ET AL: "EFFECTS OF PHOSPHOROTHIOATE CAPPING ON ANTISENSE OLIGONUCLEOTIDE STABILITY, HYBRIDIZATION AND ANTIVIRAL EFFICACY VERSUS HERPES SIMPLEX VIRUS INFECTION". see page 5747, column 1, paragraph 2 -----</p>	17-24

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 96/ 13371

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 9-16, 25-32 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compounds.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. Claims 1 - 8
2. Claims 17 - 24

1. ☒ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☒ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International Application No.

Information on patent family members

PCT/US 96/13371

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9408003 A	14-04-94	AU 5295993 A	26-04-94
		EP 0670897 A	13-09-95
		FI 951601 A	29-05-95
		HU 73335 A	29-07-96
		JP 7508657 T	28-09-95
		NO 951306 A	04-04-95
		US 5576208 A	19-11-96
		AU 672175 B	26-09-96
		AU 2249792 A	12-01-93
		BR 9206156 A	17-10-95
		CA 2111472 A	23-12-92
		EP 0590082 A	06-04-94
		JP 6504679 T	02-06-94
		WO 9222651 A	23-12-92
		US 5582986 A	10-12-96
		US 5582972 A	10-12-96
WO 9502069 A	19-01-95	AU 7398194 A	06-02-95
		BR 9406931 A	10-09-96
		CA 2166058 A	19-01-95
		EP 0714449 A	05-06-96
		FI 960089 A	07-03-96
		JP 8507929 T	27-08-96
		NO 960079 A	29-02-96
WO 9313114 A	08-07-93	AU 3249793 A	28-07-93
		CA 2126451 A	08-07-93
		EP 0644889 A	29-03-95
		JP 6511387 T	22-12-94
DE 4110085 A	01-10-92	NONE	
WO 9423028 A	13-10-94	AU 6527094 A	24-10-94
		CA 2159350 A	13-10-94
		CN 1124980 A	19-06-96
		EP 0693123 A	24-01-96
		JP 8510900 T	19-11-96
WO 9402499 A	03-02-94	AU 4688393 A	14-02-94
		CA 2140542 A	03-02-94

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 96/13371

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9402499 A		EP 0652890 A JP 8502723 T	17-05-95 26-03-96
WO 9112323 A	22-08-91	US 5149797 A AT 141326 T CA 2076037 A DE 69121391 D DE 69121391 T EP 0515511 A ES 2090314 T US 5366878 A US 5220007 A	22-09-92 15-08-96 16-08-91 19-09-96 06-03-97 02-12-92 16-10-96 22-11-94 15-06-93
WO 9407367 A	14-04-94	NONE	



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<p>(21) International Application Number: PCT/US96/13371</p> <p>(22) International Filing Date: 16 August 1996 (16.08.96)</p> <p>(30) Priority Data: 08/516,454 17 August 1995 (17.08.95) US</p> <p>(71) Applicant: HYBRIDON, INC. [US/US]; One Innovation Drive, Worcester, MA 01605 (US).</p> <p>(72) Inventor: AGRAWAL, Sudhir; 61 Lamplighter Road, Shrewsbury, MA 01545 (US).</p> <p>(74) Agent: KEOWN, Wayne, A.; Hale and Dorr, 60 State Street, Boston, MA 02109 (US).</p>		<p>(81) Designated States: AL, AU, BB, BG, BR, CA, CN, CZ, EE, GE, HU, IL, IS, JP, KP, KR, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, TR, TT, UA, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>Without international search report and to be republished upon receipt of that report.</i></p>
<p>(54) Title: INVERTED CHIMERIC AND HYBRID OLIGONUCLEOTIDES</p> <p>(57) Abstract</p> <p>The invention relates to modified oligonucleotides that are useful for studies of gene expression and for the antisense therapeutic approach. The invention provides inverted hybrid oligonucleotides and inverted chimeric oligonucleotides, both of which produce reduced side effects, relative to traditional phosphorothioate, hybrid or chimeric oligonucleotides.</p>		

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INVERTED CHIMERIC AND HYBRID OLIGONUCLEOTIDES

BACKGROUND OF THE INVENTION

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Field of the invention

The invention relates to modified oligonucleotides that are useful for studies of gene expression and for the antisense therapeutic approach.

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Summary of the related art

The potential for using oligonucleotides as inhibitors of specific gene expression in an antisense therapeutic approach was first suggested in three articles published in 15 1977 and 1978. Paterson et al., Proc. Natl. Acad. Sci. USA 74: 4370-4374 (1977) discloses that cell-free translation of mRNA can be inhibited by binding a complementary oligonucleotide to the mRNA. Zamecnik and Stephenson, Proc. Natl. Acad. Sci. USA 75: 280-284 and 285-288 (1978) disclose 20 that a 13-mer synthetic oligonucleotide that is complementary to a part of the Rous sarcoma virus (RSV) genome can inhibit RSV replication in infected cell cultures and can inhibit RSV-mediated transformation of primary chick fibroblasts into malignant sarcoma cells.

25 Since these early studies, the ability of antisense oligonucleotides to inhibit virus propagation has become firmly established. US Patent No. 4,806,463 teaches that human immunodeficiency virus propagation can be inhibited by oligonucleotides that are complementary to any of various 30 regions of the HIV genome. US Patent No. 5,194,428 discloses inhibition of influenza virus replication by phosphorothioate oligonucleotides complementary to the influenza virus polymerase 1 gene. Agrawal, Trends in Biotechnology 10: 152-158 (1992) reviews the use of 35 antisense oligonucleotides as antiviral agents.

Antisense oligonucleotides have also been developed as anti-parasitic agents. PCT publication no. WO93/13740 discloses the use of antisense oligonucleotides to inhibit propagation of drug-resistant malarial parasites. Tao et al., Antisense Research and Development 5: 123-129 (1995) teaches inhibition of propagation of a schistosome parasite by antisense oligonucleotides.

More recently, antisense oligonucleotides have shown promise as candidates for therapeutic applications for diseases resulting from expression of cellular genes. PCT publication no. WO95/09236 discloses reversal of beta amyloid-induced neuronal cell line morphological abnormalities by oligonucleotides that inhibit beta amyloid expression. PCT publication no. WO94/26887 discloses reversal of aberrant splicing of a globin gene transcript by oligonucleotides complementary to certain portions of that transcript. PCT application no. PCT/US94/13685 discloses inhibition of tumorigenicity by oligonucleotides complementary to the gene encoding DNA methyltransferase.

The development of various antisense oligonucleotides as therapeutic and diagnostic agents has recently been reviewed by Agrawal and Iyer, Current Opinion in Biotechnology 6: 12-19 (1995).

As interest in the antisense therapeutic approach has grown, various efforts have been made to improve the pharmacologic properties of oligonucleotides by modifying the sugar-phosphate backbone. US Patent No. 5,149,797 describes traditional chimeric oligonucleotides having a phosphorothioate core region interposed between methylphosphonate or phosphoramidate flanking regions. PCT publication no. WO94/02498 discloses traditional hybrid oligonucleotides having regions of 2'-O-substituted ribonucleotides flanking a DNA core region.

Much is currently being discovered about the pharmacodynamic properties of oligonucleotides. Agrawal et al., Clinical Pharmacokinetics 28: 7-16 (1995) and Zhang et

al., Clinical Pharmacology and Therapeutics 58: 44-53 (1995) disclose pharmacokinetics of anti-HIV oligonucleotides in human patients. Some of these new discoveries have led to new challenges to be overcome for the optimization of oligonucleotides as therapeutic agents. For example, Kniep et al., Nature 374: 546-549 (1995) discloses that oligonucleotides containing the CG dinucleotide flanked by certain other sequences have a mitogenic effect in vivo. Galbraith et al., Antisense Research and Development 4: 201-206 (1994) disclose complement activation by oligonucleotides. Henry et al., Pharm. Res. 11: PPDM8082 (1994) discloses that oligonucleotides may potentially interfere with blood clotting.

15 There is, therefore, a need for modified oligonucleotides that retain gene expression inhibition properties while producing fewer side effects than conventional oligonucleotides.

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BRIEF SUMMARY OF THE INVENTION

The invention relates to modified oligonucleotides that
5 are useful for studies of gene expression and for the
antisense therapeutic approach. The invention provides
modified oligonucleotides that inhibit gene expression and
that produce fewer side effects than conventional
oligonucleotides. In particular, the invention provides
10 modified oligonucleotides that demonstrate reduced
mitogenicity, reduced activation of complement and reduced
antithrombotic properties, relative to conventional
oligonucleotides.

15 In a first aspect, the invention provides inverted
hybrid and inverted chimeric oligonucleotides and
compositions of matter for inhibiting specific gene
expression with reduced side effects. Such inhibition of
gene expression can be used as an alternative to mutant
20 analysis for determining the biological function of specific
genes. Such inhibition of gene expression can also be used
to therapeutically treat diseases that are caused by
expression of the genes of a virus or a pathogen, or by the
inappropriate expression of cellular genes.

25 In one preferred embodiment according to this aspect of
the invention, the composition of matter comprises modified
oligonucleotides having one or more 2'-O-substituted RNA
region flanked by one or more oligodeoxyribonucleotide
phosphorothioate region. In certain particularly preferred
30 embodiments, the 2'-O-substituted RNA region is in between
two oligodeoxyribonucleotide regions, a structure that is
"inverted" relative to traditional hybrid oligonucleotides.
In another preferred embodiment according to this aspect of
the invention, the composition of matter comprises modified
35 oligonucleotides having one or more nonionic oligonucleotide
region flanked by one or more region of oligonucleotide

phosphorothioate. In preferred embodiments, the nonionic region contains alkylphosphonate and/or phosphoramidate and/or phosphotriester internucleoside linkages. In certain particularly preferred embodiments, the nonionic
5 oligonucleotide region is in between two oligonucleotide phosphorothioate regions, a structure that is "inverted" relative to traditional chimeric oligonucleotides.

In a second aspect, the invention provides a method for
10 modulating gene expression in a mammal with reduced side effects. In the method according to this aspect of the invention, a composition of matter according to the first aspect of the invention is administered to the mammal, wherein the oligonucleotide is complementary to a gene that
15 is being expressed in the mammal. In a preferred embodiment, after the composition of matter is administered, one or more measurement is taken of biological effects selected from the group consisting of complement activation, mitogenesis and inhibition of thrombin clot formation.

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In a third aspect, the invention provides a method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease
25 a composition of matter according to the first aspect of the invention, wherein the oligonucleotide is complementary to a gene that is aberrantly expressed, wherein such aberrant expression causes the disease. In this context, aberrant gene expression means expression in a host organism of a
30 gene required for the propagation of a virus or a prokaryotic or eukaryotic pathogen, or inappropriate expression of a host cellular gene. Inappropriate host cellular gene expression includes expression of a mutant allele of a cellular gene, or underexpression or
35 overexpression of a normal allele of a cellular gene, such that disease results from such inappropriate host cellular

gene expression. In a preferred embodiment, after the composition of matter is administered, one or more measurement is taken of biological effects selected from the group consisting of complement activation, mitogenesis and inhibition of thrombin clot formation.

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BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows inverted hybrid oligonucleotides, hybrid
5 oligonucleotides and oligonucleotide phosphodiester and
phosphorothioates used in the current studies. 2'-O-
methylribo-nucleotides are outlined and phosphodiester-
linked nucleotides are underlined; all others are
phosphorothioate-linked nucleotides.

10 Figure 2 shows mixed backbone, chimeric and inverted
chimeric oligonucleotides used in the current studies.
Methylphosphonate-linked nucleotides are underlined; all
others are phosphorothioate linked nucleotides.

Figure 3 shows thymidine uptake by mouse spleenocytes
15 as a function of concentration of phosphorothioate
oligonucleotide or any of various inverted hybrid
oligonucleotides.

Figure 4 shows extent of inhibition of complement-
mediated hemolysis observed when serum is treated with
20 phosphorothioate oligonucleotide or any of various inverted
hybrid oligonucleotides.

Figure 5 shows prolongation of aPTT obtained when
normal human serum is treated with phosphorothioate
oligonucleotides or with any of various inverted hybrid
25 oligonucleotides.

Figure 6 shows thymidine uptake by mouse spleenocytes
as a function of concentration of phosphorothioate
oligonucleotide or any of various inverted chimeric
oligonucleotides.

30 Figure 7 shows extent of inhibition of complement-
mediated hemolysis observed when serum is treated with
phosphorothioate oligonucleotide or any of various inverted
chimeric oligonucleotides.

Figure 8 shows prolongation of aPTT obtained when normal human serum is treated with phosphorothioate oligonucleotides or with any of various inverted chimeric oligonucleotides.

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DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

All US Patents, patent publications and scientific
5 literature cited in this specification evidence the level of
knowledge in the field and are hereby incorporated by
reference.

The invention relates to modified oligonucleotides that
10 are useful for studies of gene expression and for the
antisense therapeutic approach. The invention provides
modified oligonucleotides that inhibit gene expression and
that produce fewer side effects than conventional
oligonucleotides. In particular, the invention provides
15 modified oligonucleotides that demonstrate reduced
mitogenicity, reduced activation of complement and reduced
antithrombotic properties, relative to conventional
oligonucleotides.

20 In a first aspect, the invention provides inverted
hybrid and inverted chimeric oligonucleotides and
compositions of matter for inhibiting specific gene
expression with reduced side effects. Such inhibition of
gene expression can be used as an alternative to mutant
25 analysis or gene "knockout" experiments for determining the
biological function of specific genes. Such inhibition of
gene expression can also be used to therapeutically treat
diseases that are caused by expression of the genes of a
virus or a pathogen, or by the inappropriate expression of
30 cellular genes.

A composition of matter for inhibiting specific gene
expression with reduced side effects, according to this
aspect of the invention, comprises a modified
oligonucleotide that is complementary to a portion of a
35 genomic region or gene for which inhibition of expression is
desired, or to RNA transcribed from such a gene. For

purposes of the invention, the term oligonucleotide includes polymers of two or more deoxyribonucleotide, ribonucleotide, or 2'-O-substituted ribonucleotide monomers, or any combination thereof. The term oligonucleotide also
5 encompasses such polymers having chemically modified bases or sugars and/ or having additional substituents, including without limitation lipophilic groups, intercalating agents, diamines and adamantane. Preferably, such oligonucleotides will have from about 12 to about 50 nucleotides, most
10 preferably from about 17 to about 35 nucleotides. The term complementary means having the ability to hybridize to a genomic region, a gene, or an RNA transcript thereof under physiological conditions. Such hybridization is ordinarily the result of base-specific hydrogen bonding between
15 complementary strands, preferably to form Watson-Crick or Hoogsteen base pairs, although other modes of hydrogen bonding, as well as base stacking can also lead to hybridization. As a practical matter, such hybridization can be inferred from the observation of specific gene
20 expression inhibition. The gene sequence or RNA transcript sequence to which the modified oligonucleotide sequence is complementary will depend upon the biological effect that is sought to be modified. In some cases, the genomic region, gene, or RNA transcript thereof may be from a virus.
25 Preferred viruses include, without limitation, human immunodeficiency virus (type 1 or 2), influenza virus, herpes simplex virus (type 1 or 2), Epstein-Barr virus, cytomegalovirus, respiratory syncytial virus, influenza virus, hepatitis B virus, hepatitis C virus and papilloma
30 virus. In other cases, the genomic region, gene, or RNA transcript thereof may be from endogenous mammalian (including human) chromosomal DNA. Preferred examples of such genomic regions, genes or RNA transcripts thereof include, without limitation, sequences encoding vascular
35 endothelial growth factor (VEGF), beta amyloid, DNA

methyltransferase, protein kinase A, ApoE4 protein, p-glycoprotein, c-MYC protein, BCL-2 protein and CAPL. In yet other cases, the genomic region, gene, or RNA transcript thereof may be from a eukaryotic or prokaryotic pathogen including, without limitation, *Plasmodium falciparum*, *Plasmodium malarie*, *Plasmodium ovale*, *Schistosoma spp.*, and *Mycobacterium tuberculosis*.

In addition to the modified oligonucleotide according to the invention, the composition of matter for inhibiting gene expression with reduced side effects may optionally contain any of the well known pharmaceutically acceptable carriers or diluents. This composition of matter may further contain one or more additional oligonucleotides according to the invention, which additional oligonucleotide may be either an inverted hybrid oligonucleotide or an inverted chimeric oligonucleotide. Alternatively, this composition may contain one or more traditional antisense oligonucleotide, such as an oligonucleotide phosphorothioate, a hybrid oligonucleotide, or a chimeric oligonucleotide, or it may contain any other pharmacologically active agent.

In one preferred embodiment according to this aspect of the invention, the composition of matter comprises modified oligonucleotides having one or more 2'-O-substituted RNA region flanked by one or more oligodeoxyribonucleotide phosphorothioate region. In certain particularly preferred embodiments, the 2'-O-substituted RNA region is in between two oligodeoxyribonucleotide phosphorothioate regions, a structure that is "inverted" relative to traditional hybrid oligonucleotides. Accordingly, oligonucleotides according to this embodiment are designated inverted hybrid oligonucleotides. The 2'-O-substituted RNA region preferably has from about four to about 10 or 13 2'-O-substituted nucleosides joined to each other by 5' to 3' internucleoside linkages, and most preferably from about four to about eight such 2'-O-substituted nucleosides.

Preferably, the overall size of the inverted hybrid oligonucleotide will be from about 15 to about 35 or 50 nucleotides. Most preferably, the 2'-O-substituted ribonucleosides will be linked to each other through a 5' to 3' phosphorothioate, phosphotriester, or phosphodiester linkage. For purposes of the invention the term "2'-O-substituted" means substitution of the 2' position of the pentose moiety with an -O-lower alkyl group containing 1-6 saturated or unsaturated carbon atoms, or with an -O-aryl or allyl group having 2-6 carbon atoms, wherein such alkyl, aryl or allyl group may be unsubstituted or may be substituted, e.g., with halo, hydroxy, trifluoromethyl, cyano, nitro, acyl, acyloxy, alkoxy, carboxyl, carbalkoxyl, or amino groups; or with a hydroxy, an amino or a halo group, but not with a 2'-H group. The phosphorothioate flanking region or regions has from about four to about 46 nucleosides joined to each other by 5' to 3' phosphorothioate linkages, and preferably from about 5 to about 26 such phosphorothioate-linked nucleosides. Most preferably, the phosphorothioate regions will have from about 5 to about 15 phosphorothioate-linked nucleosides. The phosphorothioate linkages may be mixed R_p and S_p enantiomers, or they may be stereoregular or substantially stereoregular in either R_p or S_p form (see Iyer et al., Tetrahedron Asymmetry 6: 1051-1054 (1995)).

In another preferred embodiment according to this aspect of the invention, the composition of matter comprises modified oligonucleotides having one or more nonionic oligonucleotide region flanked by one or more region of oligonucleotide phosphorothioate. In preferred embodiments, the nonionic region contains alkylphosphonate and/or phosphoramidate and/or phosphotriester internucleoside linkages. In certain particularly preferred embodiments, the nonionic oligonucleotide region is in between two oligonucleotide phosphorothioate regions, a structure that is "inverted" relative to traditional chimeric

oligonucleotides. Accordingly, oligonucleotides according to this embodiment are designated inverted chimeric oligonucleotides. The nonionic region has from about four to about 10 or 12 nucleosides joined to each other by 5' to 3' nonionic linkages, preferably alkylphosphonate, phosphoramidate or phosphotriester linkages, and preferably from about four to about eight such nonionic-linked nucleosides. The phosphorothioate flanking region or regions has from about four to about 46 nucleosides joined to each other by 5' to 3' phosphorothioate linkages, and preferably from about eight to about 26 such phosphorothioate-linked nucleosides. Most preferably, the phosphorothioate regions will have from about 5 to about 15 phosphorothioate-linked nucleosides. The phosphorothioate linkages may be mixed R_p and S_p enantiomers, or they may be stereoregular or substantially stereoregular in either R_p or S_p form (see Iyer et al., Tetrahedron Asymmetry 6: 1051-1054 (1995)). In a most preferred embodiment the oligonucleotide has a nonionic region having from about 6 to about 8 methylphosphonate-linked nucleosides, flanked on either side by phosphorothioate regions, each having from about 6 to about 10 phosphorothioate-linked nucleosides.

Those skilled in the art will recognize that the elements of these preferred embodiments can be combined and the inventor does contemplate such combination. For example, 2'-O-substituted ribonucleotide regions may well include from one to all nonionic internucleoside linkages. Alternatively, nonionic regions may have from one to all 2'-O-substituted ribonucleotides. Moreover, oligonucleotides according to the invention may contain combinations of one or more 2'-O-substituted ribonucleotide region and one or more nonionic region, either or both being flanked by phosphorothioate regions. (See Nucleosides & Nucleotides 14: 1031-1035 (1995) for relevant synthetic techniques.

In a second aspect, the invention provides a method for modulating gene expression in a mammal with reduced side effects. In the method according to this aspect of the invention, a composition of matter according to the first
5 aspect of the invention is administered to the mammal, wherein the oligonucleotide is complementary to a gene that is being expressed in the mammal. Preferably, such administration may be parenteral, oral, intranasal or intrarectal. In a preferred embodiment, after the
10 composition of matter is administered, one or more measurement is taken of biological side effects selected from the group consisting of complement activation, mitogenesis and inhibition of thrombin clot formation.

15 In a third aspect, the invention provides a method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to the first aspect of the
20 invention, wherein the oligonucleotide is complementary to a gene that is aberrantly expressed, wherein such aberrant expression causes the disease. In this context, aberrant gene expression means expression in a host organism of a gene required for the propagation of a virus or a
25 prokaryotic or eukaryotic pathogen, or inappropriate expression of a host cellular gene. Inappropriate host cellular gene expression includes expression of a mutant allele of a cellular gene, or underexpression or
30 overexpression of a normal allele of a cellular gene, such that disease results from such inappropriate host cellular gene expression. Preferably, such administration should be parenteral, oral, sublingual, transdermal, topical, intranasal or intrarectal. Administration of the therapeutic
35 compositions can be carried out using known procedures at dosages and for periods of time effective to reduce symptoms or surrogate markers of the disease. When administered

systemically, the therapeutic composition is preferably administered at a sufficient dosage to attain a blood level of oligonucleotide from about 0.01 micromolar to about 10 micromolar. For localized administration, much lower concentrations than this may be effective, and much higher concentrations may be tolerated. Preferably, a total dosage of oligonucleotide will range from about 0.1 mg oligonucleotide per patient per day to about 200 mg oligonucleotide per kg body weight per day. It may desirable to administer simultaneously, or sequentially a therapeutically effective amount of one or more of the therapeutic compositions of the invention to an individual as a single treatment episode. In a preferred embodiment, after the composition of matter is administered, one or more measurement is taken of biological effects selected from the group consisting of complement activation, mitogenesis and inhibition of thrombin clot formation.

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The following examples are intended to further illustrate certain preferred embodiments of the invention and are not intended to limit the scope of the invention.

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Example 1

Synthesis, Deprotection And Purification Of Oligonucleotides

Oligonucleotide phosphorothioates were synthesized using an automated DNA synthesizer (Model 8700, Biosearch, Bedford, MA) using a beta-cyanoethyl phosphoramidite approach on a 10 micromole scale. To generate the phosphorothioate linkages, the intermediate phosphite linkage obtained after each coupling was oxidized using 3H, 1,2-benzodithiole-3H-one-1,1-dioxide (See Beaucage, In *Protocols for Oligonucleotides and Analogs: Synthesis and Properties*, Agrawal (editor), Humana Press, Totowa, NJ, pp. 33-62 (1993).) Similar synthesis was carried out to generate phosphodiester linkages, except that a standard oxidation was carried out using standard iodine reagent. Synthesis of inverted chimeric oligonucleotide was carried out in the same manner, except that methylphosphonate linkages were assembled using nucleoside methylphosphonamidite (Glen Research, Sterling, VA), followed by oxidation with 0.1 M iodine in tetrahydrofuran/2,6-lutidine/water (75:25:0.25) (see Agrawal & Goodchild, *Tet. Lett.* 28: 3539-3542 (1987)). Inverted hybrid oligonucleotides were synthesized similarly, except that the segment containing 2'-O-methylribonucleotides was assembled using 2'-O-methylribonucleoside phosphoramidite, followed by oxidation to a phosphorothioate or phosphodiester linkage as described above. Deprotection and purification of oligonucleotides was carried out according to standard procedures, (See Padmapriya et al., *Antisense Res. & Dev.* 4: 185-199 (1994)),

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except for oligonucleotides containing methylphosphonate-containing regions. For those oligonucleotides, the CPG-bound oligonucleotide was treated with concentrated ammonium hydroxide for 1 hour at room temperature, and the
5 supernatant was removed and evaporated to obtain a pale yellow residue, which was then treated with a mixture of ethylenediamine/ethanol (1:1 v/v) for 6 hours at room temperature and dried again under reduced pressure.

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Example 2

Reduced Complement Activation In Vitro By Inverted Hybrid And Inverted Chimeric Oligonucleotides

To determine the relative effect of inverted hybrid or
15 inverted chimeric structure on oligonucleotide-mediated depletion of complement, the following experiments were performed. Venous blood was collected from healthy adult human volunteers. Serum was prepared for hemolytic complement assay by collecting blood into vacutainers
20 (Becton Dickinson #6430 Franklin Lakes, NJ) without commercial additives. Blood was allowed to clot at room temperature for 30 minutes, chilled on ice for 15 minutes, then centrifuged at 4°C to separate serum. Harvested serum was kept on ice for same day assay or, alternatively, stored
25 at -70°C. Buffer, oligonucleotide phosphorothioate, inverted hybrid oligonucleotide, or inverted chimeric oligonucleotide was then incubated with the serum. A standard CH50 assay (see Kabat and Mayer (eds): *Experimental Immunochimistry*, 2d Edition, Springfield, IL, CC Thomas
30 (1961), p.125) for complement-mediated lysis of sheep red blood cells (Colorado Serum Co.) sensitized with anti-sheep red cell antibody (hemolysin, Diamedix, Miami, FL) was performed, using duplicate determinations of at least five dilutions of each test serum, then hemoglobin release into
35 cell-free supernates was measured spectrophotometrically at 541 nm.

Example 3**Reduced Mitogenicity In Vitro Of Inverted Hybrid
And Inverted Chimeric Oligonucleotides**

5 To determine the relative effect of inverted hybrid or inverted chimeric structure on oligonucleotide-mediated mitogenicity, the following experiments were performed. Spleen was taken from a male CD1 mouse (4-5 weeks, 20-22 g; Charles River, Wilmington, MA). Single cell suspensions
10 were prepared by gently mincing with frosted edges of glass slides. Cells were then cultured in RPMI complete media [RPMI media supplemented with 10% fetal bovine serum (FBS), 50 micromolar 2-mercaptoethanol (2-ME), 100 U/ml penicillin, 100 micrograms/ml streptomycin, 2 mM L-glutamine]. To
15 minimize oligonucleotide degradation, FBS was first heated for 30 minutes at 65°C (phosphodiester-containing oligonucleotides) or 56°C (all other oligonucleotides). Cells were plated in 96 well dishes at 100,000 cells per well (volume of 100 microliters/ well). Oligonucleotides in
20 10 microliters TE buffer (10mM Tris-HCl, pH 7.5, 1mM EDTA) were added to each well. After 44 hours of culturing at 37°C, one microcurie tritiated thymidine (Amersham, Arlington Heights, IL) was added in 20 microliters RPMI media for a 4 hour pulse labelling. The cells were then
25 harvested in an automatic cell harvester (Skatron, Sterling, VA) and the filters were assessed using a scintillation counter. In control experiments for mitogenicity, cells were treated identically, except that either media (negative control) or concanavalin A (positive control) was added to
30 the cells in place of the oligonucleotides. The results of these studies are shown in Figure 1. All of the inverted hybrid oligonucleotides proved to be less immunogenic than phosphorothioate oligonucleotides. Inverted hybrid oligonucleotides having phosphodiester linkages in the 2'-O-
35 methyl region appeared to be slightly less immunogenic than those containing phosphorothioate linkages in that region.

No significant difference in mitogenicity was observed when the 2'-O-methylribonucleotide region was pared down from 13 to 11 or to 9 nucleotides. Inverted chimeric oligonucleotides were also generally less mitogenic than phosphorothioate oligonucleotides. In addition, these oligonucleotides appeared to be less mitogenic than traditional chimeric oligonucleotides, at least in cases in which the traditional chimeric oligonucleotides had significant numbers of methylphosphonate linkages near the 3' end. Increasing the number of methylphosphonate linkers in the middle of the oligonucleotide from 5 to 6 or 7 did not appear to have a significant effect on mitogenicity. These results indicate that incorporation of inverted hybrid or inverted chimeric structure into an oligonucleotide can reduce its mitogenicity.

Example 4

Reduced Inhibition Of Clotting In Vitro By Inverted Hybrid And Inverted Chimeric Oligonucleotides

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To determine the relative effect of inverted hybrid or inverted chimeric structure on oligonucleotide-induced mitogenicity, the following experiments were performed. Venous blood was collected from healthy adult human volunteers. Plasma for clotting time assay was prepared by collecting blood into siliconized vacutainers with sodium citrate (Becton Dickinson #367705), followed by two centrifugations at 4°C to prepare platelet-poor plasma. Plasma aliquots were kept on ice, spiked with various test compounds, and either tested immediately or quickly frozen on dry ice for subsequent storage at -20°C prior to coagulation assay. Activated partial thromboplastin time (aPTT) was performed in duplicate on an Electra 1000C (Medical Laboratory Automation, Mount Vernon, NY) according to the manufacturer's recommended procedures, using Actin FSL (Baxter Dade, Miami, FL) and calcium to initiate clot

formation, which was measured photometrically. Prolongation of aPTT was taken as an indication of clotting inhibition side effect produced by the oligonucleotide. The results are shown in Figure 5 for inverted hybrid oligonucleotides and in Figure 8 for inverted chimeric oligonucleotides. Traditional phosphorothioate oligonucleotides produce the greatest prolongation of aPTT, of all of the oligonucleotides tested. Traditional hybrid oligonucleotides produced somewhat reduced prolongation of aPTT. In comparison with traditional phosphorothioate or traditional hybrid oligonucleotides, all of the inverted hybrid oligonucleotides tested produced significantly reduced prolongation of aPTT. Inverted hybrid oligonucleotides having phosphodiester linkages in the 2'-O-substituted ribonucleotide region had the greatest reduction in this side effect, with one such oligonucleotide having a 2'-O-methyl RNA phosphodiester region of 13 nucleotides showing very little prolongation of aPTT, even at oligonucleotide concentrations as high as 100 micrograms/ml. Traditional chimeric oligonucleotides produce much less prolongation of aPTT than do traditional phosphorothioate oligonucleotides. Generally, inverted chimeric oligonucleotides retain this characteristic. At least one inverted chimeric oligonucleotide, having a methylphosphonate region of 7 nucleotides flanked by phosphorothioate regions of 9 nucleotides, gave better results in this assay than the traditional chimeric oligonucleotides at all but the highest oligonucleotide concentrations tested. These results indicate that inverted hybrid and inverted chimeric oligonucleotides may provide advantages in reducing the side effect of clotting inhibition when they are administered to modulate gene expression *in vivo*.

Example 5**Reduced Complement Activation In Vivo By Inverted
Hybrid And Inverted Chimeric Oligonucleotides**

5 Rhesus monkeys (4-9 kg body weight) are acclimatized to laboratory conditions for at least 7 days prior to the study. On the day of the study, each animal is lightly sedated with ketamine-HCl (10 mg/kg) and diazepam (0.5 mg/
10 kg). Surgical level anesthesia is induced and maintained by continuous ketamine intravenous drip throughout the procedure. Phosphorothioate oligonucleotide or inverted hybrid or inverted chimeric oligonucleotide is dissolved in normal saline and infused intravenously via a cephalic vein
15 catheter, using a programmable infusion pump at a delivery rate of 0.42 ml/ minute. For each oligonucleotide, oligonucleotide doses of 0, 0.5, 1, 2, 5 and 10 mg/ kg are administered to two animals each over a 10 minute infusion period. Arterial blood samples are collected 10 minutes
20 prior to oligonucleotide administration and 2, 5, 10, 20, 40 and 60 minutes after the start of the infusion, as well as 24 hours later. Serum is used for determining complement CH50, using the conventional complement-dependent lysis of sheep erythrocyte procedure (see Kabat and Mayer, 1961,
25 supra). At the highest dose, phosphorothioate oligonucleotide causes a decrease in serum complement CH50 beginning within 5 minutes of the start of infusion. Inverted hybrid and chimeric oligonucleotides are expected to show a much reduced or undetectable decrease in serum
30 complement CH50 under these conditions.

Example 6**Reduced Mitogenicity In Vivo Of Inverted Hybrid
And Inverted Chimeric Oligonucleotides**

5 CD1 mice are injected intraperitoneally with a dose of
50 mg/kg body weight of phosphorothioate oligonucleotide,
inverted hybrid oligonucleotide or inverted chimeric
oligonucleotide. Forty-eight hours later, the animals are
euthanized and the spleens are removed and weighed. Animals
10 treated with inverted hybrid or inverted hybrid
oligonucleotides are expected to show no significant
increase in spleen weight, while those treated with
oligonucleotide phosphorothioates are expected to show
modest increases in spleen weight.

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Example 7**Reduced Inhibition Of Clotting In Vivo By Inverted
Hybrid And Inverted Chimeric Oligonucleotides**

20 Rhesus monkeys are treated as in example 5. From the
whole blood samples taken, plasma for clotting assay is
prepared, and the assay performed, as described in example
4. It is expected that prolongation of aPTT will be
substantially reduced for both inverted hybrid
25 oligonucleotides and for inverted chimeric oligonucleotides,
relative to traditional oligonucleotide phosphorothioates.

Example 8**Effect Of Inverted Hybrid Or Chimeric
Structure On RNase H Activity**

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To determine the ability of inverted hybrid
oligonucleotides and inverted chimeric oligonucleotides to
35 activate RNase H when bound to a complementary RNA molecule,
the following experiments were performed. Each

oligonucleotide phosphorothioate, inverted hybrid oligonucleotide or inverted chimeric oligonucleotide was incubated together with a molar equivalent quantity of complimentary oligoribonucleotide (0.266 micromolar concentration of each), in a cuvette containing a final volume of 1 ml RNase H buffer (20 mM Tris-HCl, pH 7.5, 10 mM $MgCl_2$, 0.1 M KCl, 2% glycerol, 0.1 mM DTT). The samples were heated to 95°C, then cooled gradually to room temperature to allow annealing to form duplexes. Annealed duplexes were incubated for 10 minutes at 37°C, then 5 units RNase H was added and data collection commenced over a three hour period. Data was collected using a GBC 920 (GBC Scientific Equipment, Victoria, Australia) spectrophotometer at 259 nm. RNase H degradation was determined by hyperchromic shift.

The results are shown in Table I, below.

TABLE I

RNase H Degradation Of Oligonucleotides

<u>Oligo No. (Features)</u>	<u>Half-Life</u>	<u>Oligo No. (Features)</u>	<u>Half-Life</u>
20 GEM91 (all PO)	8.8 sec.	Hyb115 (5' MP)	11.5 sec.
GEM91 (all PS)	22.4 sec.	Hyb116 (chimeric)	9.7 sec.
GEM91H (hybrid)	32.7 sec.	Hyb117 (chimeric)	8.1 sec.
Hyb108 (inv. hyb.)	15.4 sec.	Hyb118 (inv. chim.)	11.5 sec.
Hyb109 (inv. hyb.)	7.9 sec.	Hyb119 (inv. chim.)	14.4 sec.
25 Hyb110 (inv. hyb.)	10.4 sec.	Hyb120 (inv. chim.)	9.3 sec.
Hyb111 (inv. hyb.)	12.9 sec.	Hyb121 (3' MP)	21.2 sec.
Hyb112 (inv. hyb.)	12.5 sec.	Hyb122 (chimeric)	23.0 sec.
Hyb113 (inv. hyb.)	10.9 sec.	Hyb123 (chimeric)	41.8 sec.
Hyb114 (inv. hyb.)	20.3 sec.	Hyb124 (chimeric)	not
30 detect.			

As expected, phosphodiester oligonucleotides behaved as very good co-substrates for RNase H-mediated degradation of RNA, with a degradative half-life of 8.8 seconds. Phosphorothioate oligonucleotides produced an increased half-life of 22.4 seconds. Introduction of a 2'-O-methylribonucleotide segment at either end of the

oligonucleotide further worsened RNaseH activity (half-life= 32.7 seconds). In contrast, introducing a 2'-O-methyl segment into the middle of the oligonucleotide (inverted hybrid structure) always resulted in improved RNase H-mediated degradation. When a region of 13 2'-O-methylribonucleoside phosphodiester was flanked on both sides by phosphorothioate DNA, the best RNase H activity was observed, with a half-life of 7.9 seconds. Introduction of large blocks of methylphosphonate-linked nucleosides at the 3' end of the oligonucleotide either had no effect or caused further deterioration of RNase H activity even when in a chimeric configuration. Introduction of methylphosphonate linked nucleosides at the 5' end, however, improved RNase H activity, particularly in a chimeric configuration with a single methylphosphonate linker at the 3' end (best half-life = 8.1 seconds). All inverted chimeric oligonucleotides with methylphosphonate core regions flanked by phosphorothioate regions gave good RNase results, with a half-life range of 9.3 to 14.4 seconds. These results indicate that the introduction of inverted hybrid or inverted chimeric structure into phosphorothioate-containing oligonucleotides can restore some or all of the ability of the oligonucleotide to act as a co-substrate for RNase H, a potentially important attribute for an effective antisense agent.

Example 9**Effect Of Inverted Hybrid Or Chimeric
Structure On Melting Temperature**

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To determine the effect of inverted hybrid or inverted chimeric structure on stability of the duplex formed between an antisense oligonucleotide and a target molecule, the following experiments were performed. Thermal melting (T_m) data were collected using a GBC 920 spectrophotometer, which has six 10 mm cuvettes mounted in a dual carousel. In the T_m experiments, the temperature was directed and controlled through a peltier effect temperature controller by a computer, using software provided by GBC, according to the manufacturer's directions. T_m data were analyzed by both the first derivative method and the mid-point method, as performed by the software. T_m experiments were performed in a buffer containing 10 mM PIPES, pH 7.0, 1 mM EDTA, 1 M NaCl. A VWR 1166 (VWR, Boston, MA) refrigerated bath was connected to the peltier-effect temperature controller to absorb the heat. Oligonucleotide strand concentration was determined using absorbance values at 260 nm, taking into account extinction coefficients.

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The results are shown in Table II, below.

TABLE II

Duplex Stability Of Oligonucleotides

<u>Oligo No. (Features)</u>		<u>Tm (°C) *</u>	<u>Oligo No. (Features)</u>		<u>Tm (°C) *</u>
5	GEM91 (all PO)	72.0	Hyb115 (5' MP)		61.8
	GEM91 (all PS)	63.6	Hyb116 (chimeric)		61.0
	GEM91H (hybrid)	67.0	Hyb117 (chimeric)		60.5
	Hyb108 (inv. hyb.)	76.4	Hyb118 (inv. chim.)		57.9
	Hyb109 (inv. hyb.)	80.0	Hyb119 (inv. chim.)		57.7
10	Hyb110 (inv. hyb.)	74.2	Hyb120 (inv. chim.)		56.8
	Hyb111 (inv. hyb.)	76.9	Hyb121 (3' MP)		60.7
	Hyb112 (inv. hyb.)	72.1	Hyb122 (chimeric)		60.5
	Hyb113 (inv. hyb.)	74.3	Hyb123 (chimeric)		59.0
	Hyb114 (inv. hyb.)	71.3	Hyb124 (chimeric)		not
15	detect.				

*=with complementary RNA

These results demonstrate that conversion of a phosphodiester oligonucleotide to a phosphorothioate oligonucleotide results in a reduction of duplex stability, and that introduction of methylphosphonate linkages further reduces duplex stability. Duplex stability can be restored by adding 2'-O-methylribonucleotides, and can exceed that of the phosphodiester oligonucleotide when an inverted hybrid structure is used. Conversely, use of an inverted chimeric structure results in the lowest melting temperatures observed for any hybridizing

methyolphosphonate-containing oligonucleotide, although duplex stability was still well above physiological temperatures. Taken together, these results suggest that inverted hybrid or inverted chimeric structure can be used to custom design oligonucleotides for particular duplex stabilities desired in particular experimental or therapeutic applications.

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What is claimed is:

1. An inverted hybrid oligonucleotide comprising a region
5 of 2'-O-substituted ribonucleotides in between two
oligodeoxyribonucleo-tide phosphorothioate regions.
2. The inverted hybrid oligonucleotide according to claim
1, wherein the oligonucleotide has from about 15 to about 50
nucleotides.
- 10 3. The inverted hybrid oligonucleotide according to claim
2, wherein the region of 2'-O-substituted ribonucleotides
has from about 4 to about 13 nucleotides.
4. The inverted hybrid oligonucleotide according to claim
3, wherein the region of 2'-O-substituted ribonucleotides
15 has from about 4 to about 10 nucleotides.
5. A composition of matter for inhibiting gene expression
with reduced side effects, the composition comprising the
inverted hybrid oligonucleotide according to claim 1.
6. A composition of matter for inhibiting gene expression
20 with reduced side effects, the composition comprising the
inverted hybrid oligonucleotide according to claim 2.
7. A composition of matter for inhibiting gene expression
with reduced side effects, the composition comprising the
inverted hybrid oligonucleotide according to claim 3.
- 25 8. A composition of matter for inhibiting gene expression
with reduced side effects, the composition comprising the
inverted hybrid oligonucleotide according to claim 4.
9. A method for modulating gene expression in a mammal
with reduced side effects, the method comprising the step of
30 administering to the mammal a composition of matter
according to claim 5, wherein the oligonucleotide is
complementary to a gene that is being expressed in the
mammal.

10. A method for modulating gene expression in a mammal with reduced side effects, the method comprising the step of administering to the mammal a composition of matter according to claim 6, wherein the oligonucleotide is
5 complementary to a gene that is being expressed in the mammal.

11. A method for modulating gene expression in a mammal with reduced side effects, the method comprising the step of administering to the mammal a composition of matter
10 according to claim 7, wherein the oligonucleotide is complementary to a gene that is being expressed in the mammal.

12. A method for modulating gene expression in a mammal with reduced side effects, the method comprising the step of
15 administering to the mammal a composition of matter according to claim 8, wherein the oligonucleotide is complementary to a gene that is being expressed in the mammal.

13. A method for therapeutically treating, with reduced
20 side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to claim 5, wherein the oligonucleotide is complementary to a gene that is aberrantly expressed.

25 14. A method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to claim 6, wherein the oligonucleotide is complementary to a gene that
30 is aberrantly expressed.

15. A method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to claim 7,
35 wherein the oligonucleotide is complementary to a gene that is aberrantly expressed.

16. A method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to claim 8, wherein the oligonucleotide is complementary to a gene that is aberrantly expressed.
17. An inverted chimeric oligonucleotide comprising an oligonucleotide nonionic region in between two oligonucleotide phosphorothioate regions.
18. The inverted chimeric oligonucleotide according to claim 17, wherein the oligonucleotide has from about 12 to about 50 nucleotides.
19. The inverted chimeric oligonucleotide according to claim 18, wherein the oligonucleotide nonionic region is an oligonucleotide alkylphosphonate region having from about 4 to about 12 nucleotides.
20. The inverted chimeric oligonucleotide according to claim 19, wherein the oligonucleotide alkylphosphonate region has from about 4 to about 10 nucleotides.
21. A composition of matter for inhibiting gene expression with reduced side effects, the composition comprising the inverted chimeric oligonucleotide according to claim 17.
22. A composition of matter for inhibiting gene expression with reduced side effects, the composition comprising the inverted hybrid oligonucleotide according to claim 18.
23. A composition of matter for inhibiting gene expression with reduced side effects, the composition comprising the inverted hybrid oligonucleotide according to claim 19.
24. A composition of matter for inhibiting gene expression with reduced side effects, the composition comprising the inverted hybrid oligonucleotide according to claim 20.

25. A method for modulating gene expression in a mammal with reduced side effects, the method comprising the step of administering to the mammal a composition of matter according to claim 21, wherein the oligonucleotide is
5 complementary to a gene that is being expressed in the mammal.

26. A method for modulating gene expression in a mammal with reduced side effects, the method comprising the step of administering to the mammal a composition of matter
10 according to claim 22, wherein the oligonucleotide is complementary to a gene that is being expressed in the mammal.

27. A method for modulating gene expression in a mammal with reduced side effects, the method comprising the step of
15 administering to the mammal a composition of matter according to claim 23, wherein the oligonucleotide is complementary to a gene that is being expressed in the mammal.

28. A method for modulating gene expression in a mammal
20 with reduced side effects, the method comprising the step of administering to the mammal a composition of matter according to claim 24, wherein the oligonucleotide is complementary to a gene that is being expressed in the mammal.

25 29. A method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to claim 21, wherein the oligonucleotide is complementary to a gene that
30 is aberrantly expressed.

30. A method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to claim 22,
35 wherein the oligonucleotide is complementary to a gene that is aberrantly expressed.

31. A method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having the disease a composition of matter according to claim 23,
5 wherein the oligonucleotide is complementary to a gene that is aberrantly expressed.

32. A method for therapeutically treating, with reduced side effects, a disease caused by aberrant gene expression, the method comprising administering to an individual having
10 the disease a composition of matter according to claim 24, wherein the oligonucleotide is complementary to a gene that is aberrantly expressed.

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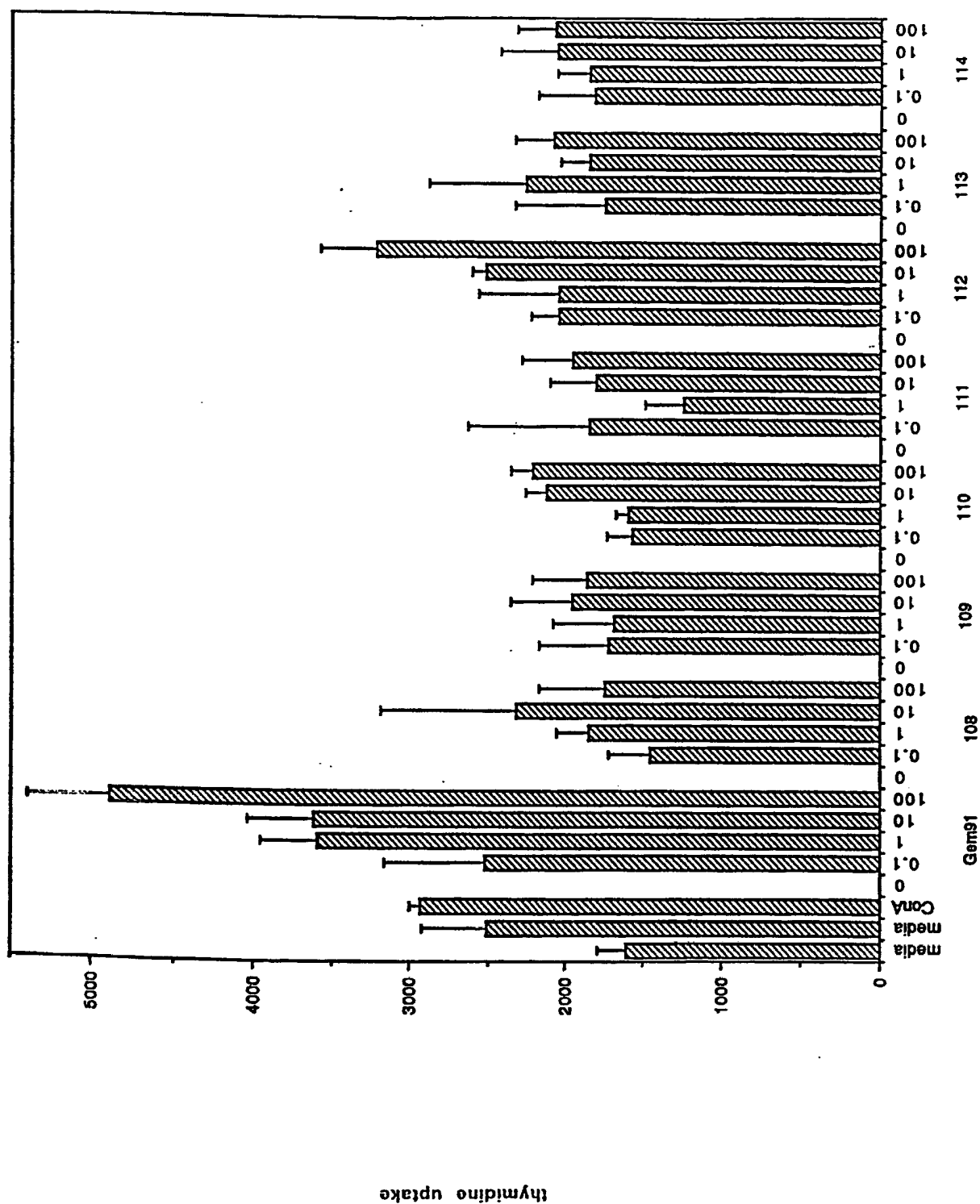
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Hyb109 5' CTC TCG CAC CCA UCU CUC UCC TTC T [SEQ ID NO:2]
Hyb110 5' CTC TCG CAC CCA UCU CUC TCC TTC T [SEQ ID NO:3]
Hyb111 5' CTC TCG CAC CCA UCU CUC TCC TTC T [SEQ ID NO:4]
Hyb112 5' CTC TCG CAC CCA UCU CUC TCC TTC T [SEQ ID NO:5]
Hyb113 5' CTC TCG CAC CCA UCU CUC TCC TTC T [SEQ ID NO:6]
Hyb114 5' CTC TCG CAC CCA UCU CTC TCC TTC T [SEQ ID NO:7]
Gem91-H 5' CUC UCG CAC CCA TCT CTC TCC UUC U [SEQ ID NO:8]
Gem91 5' CTC TCG CAC CCA TCT CTC TCC TTC T [SEQ ID NO:9]
Gem91PO 5' CTC TCG CAC CCA TCT CTC TCC TTC T [SEQ ID NO:10]

Fig. 1

- Hyb115 5' CTCTCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:11]
Hyb116 5' CTCTCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:12]
Hyb117 5' CTCTCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:13]
Hyb118 5' CTC TCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:14]
Hyb119 5' CTC TCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:15]
Hyb120 5' CTC TCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:16]
Hyb121 5' CTC TCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:17]
Hyb122 5' CTCTCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:18]
Hyb123 5' CTCTCG CAC CCA TCT CTC TCC TTC T 3' [SEQ ID NO:19]
Hyb124 5' CTT CCT CTC TCT ACC ACA GCT CTC T 3' [SEQ ID NO:20]

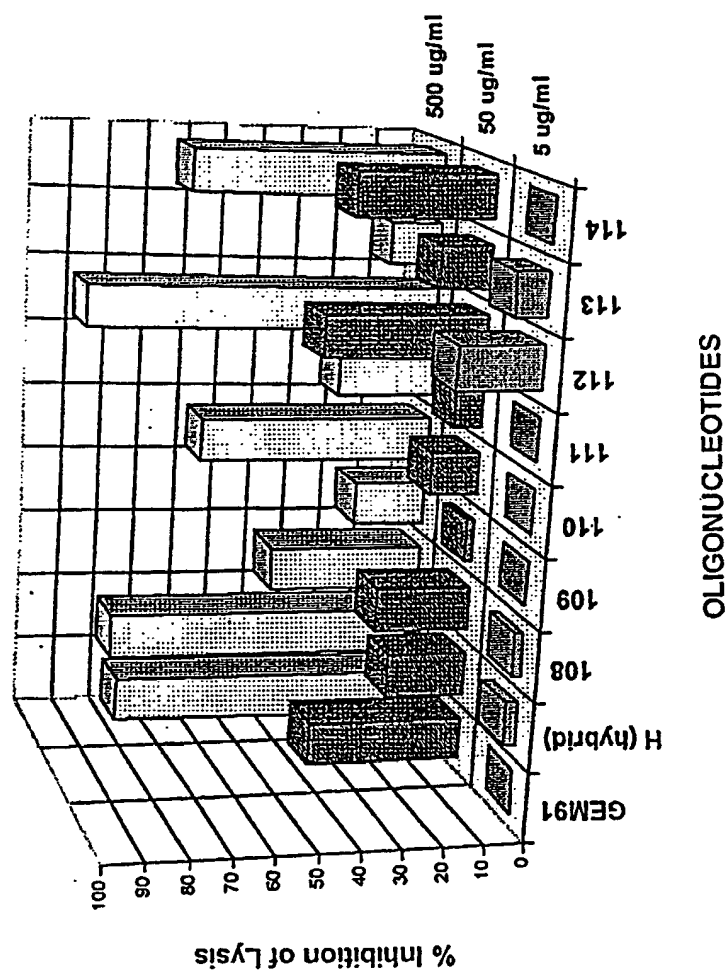
Fig. 2

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Fig. 4



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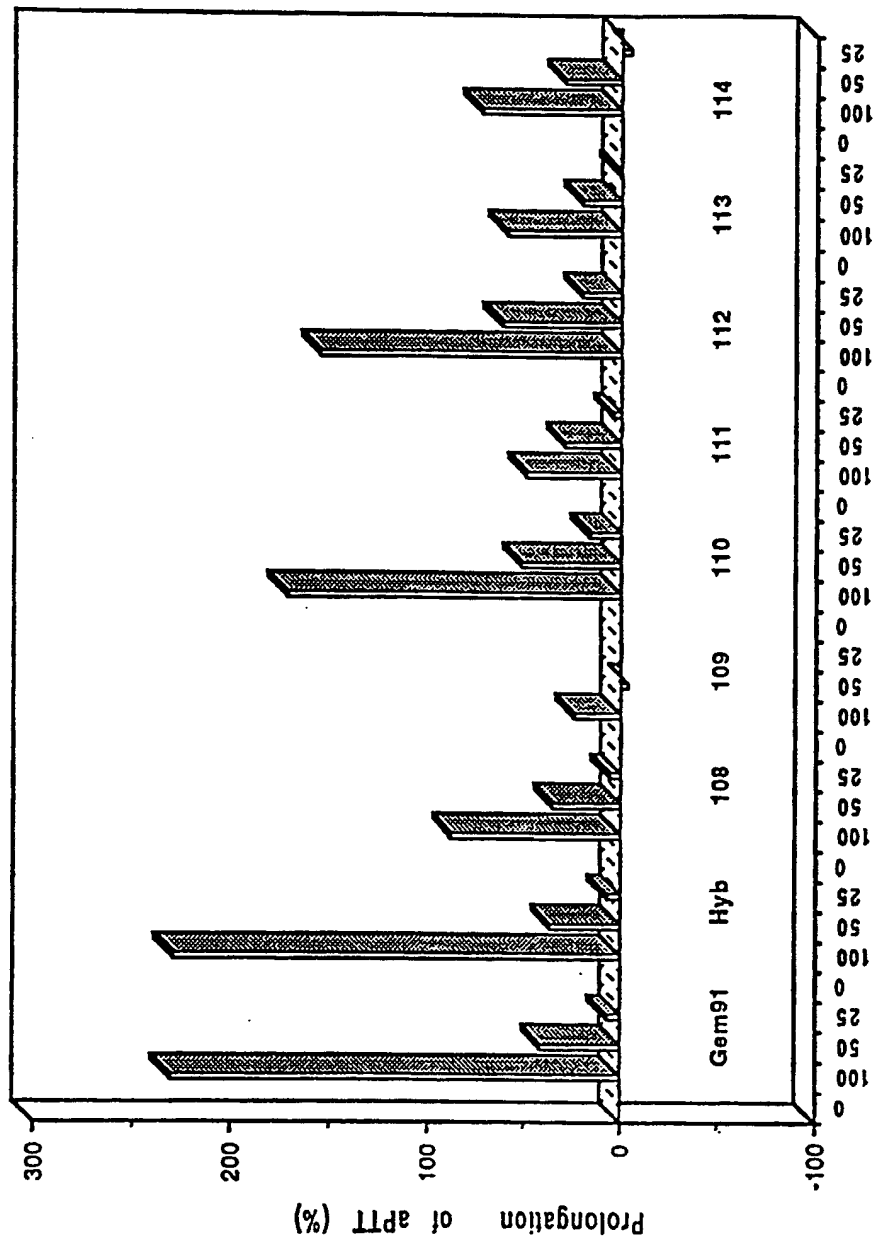
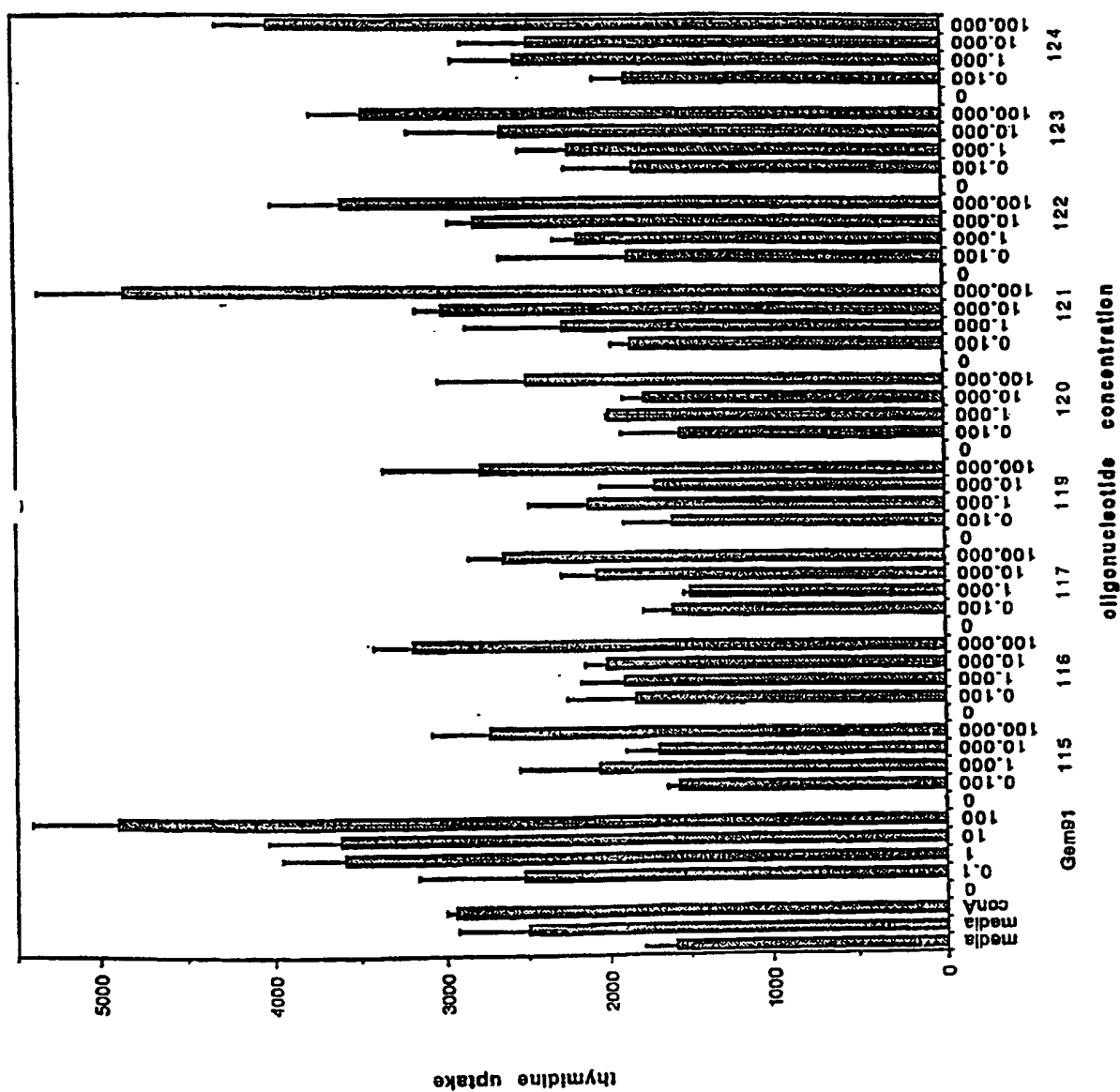


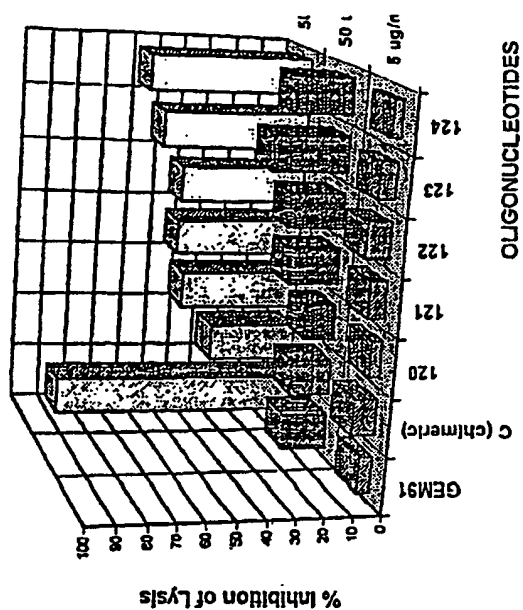
Fig 5

Concentration (µg/ml)

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Complement Data with compounds HYB0108 - 0124

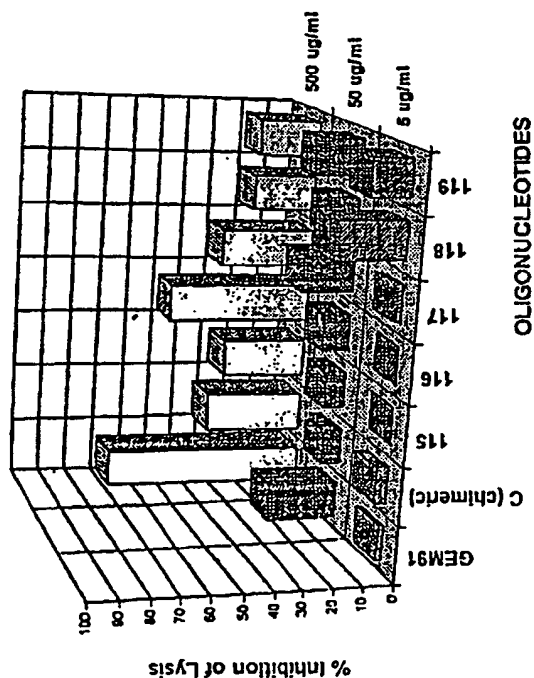


Fig. 7

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Fig 8

